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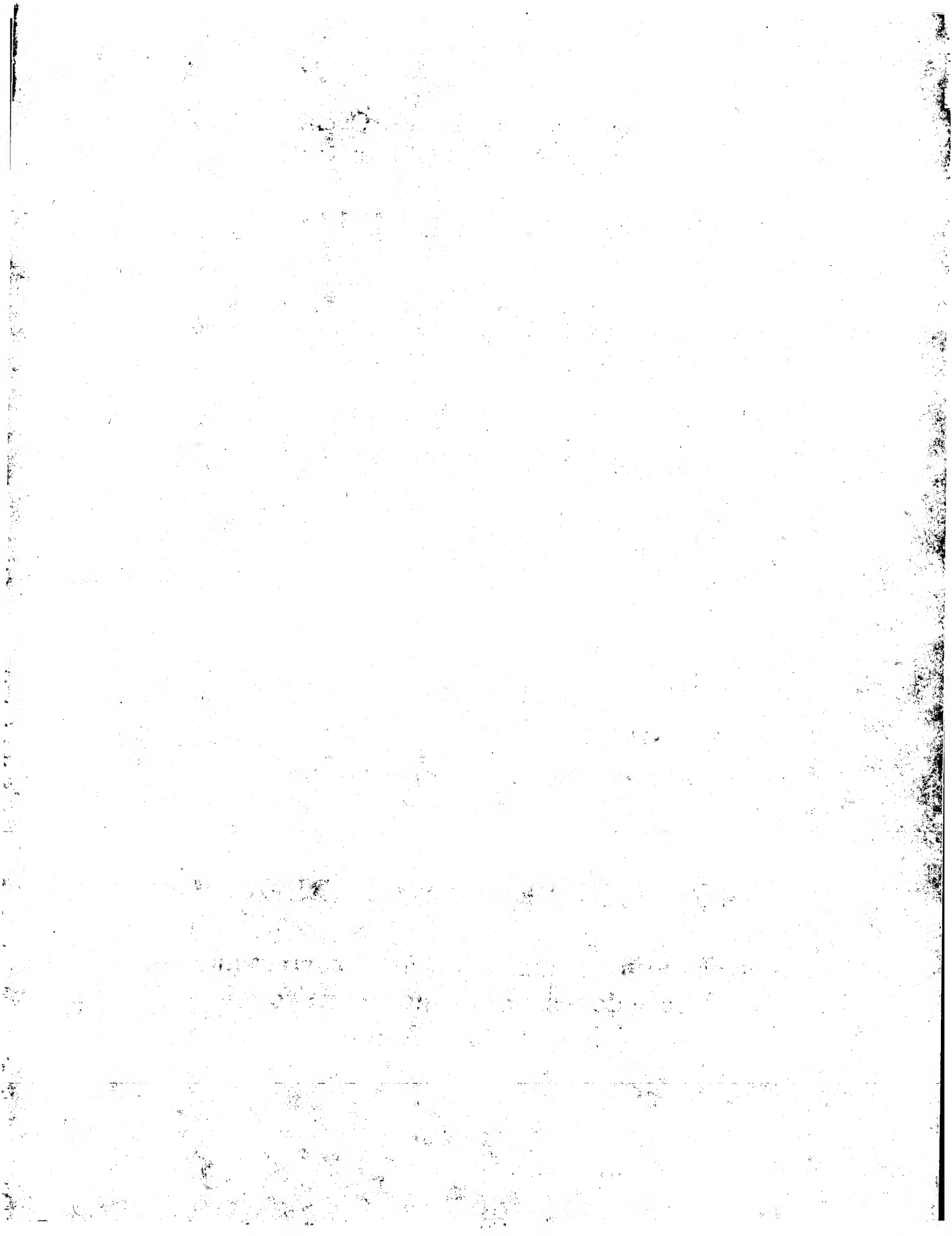
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Description

The invention relates to free-standing water-soluble polymeric films and more particularly to such films in the form of pouches and containing alkaline or borate-containing cleaning compositions.

5 A great deal of art relates to water-soluble polymeric films including polyvinyl alcohol. Much of the art has been addressed to the problem of packaging materials in such water-soluble films. As used herein, the term film describes a continuous, homogenous, dimensionally stable polymer having a small thickness in relation to area. As also used herein "polymer" means a macromolecule made up of a plurality of chemical subunits (monomers). The monomers may be identical or chemically similar, or may be of several different

10 types. Unless a more specific term is used, "polymer" will be taken to include hetero- and homo-polymers, and random, alternating, block and graft copolymers. "Copolymer" will be used to specifically refer to those polymers made from two different repeating chemical monomers. An effective water-soluble package would simplify dispensing, dispersing, slurrying, or dissolving materials contained within, as the entire package could be dumped into a mixing vessel without the need to pour out the contents. Water-soluble film packages could be used where the contents are toxic or messy, where the contents must be accurately measured, or maintained in an isolated environment, and further allow delivery of materials which are only metastable when combined, and which would ordinarily separate during storage. Soluble pre-measured

15 pouches aid convenience of consumer use in a variety of applications, particularly those involving cleaning compositions. Such cleaning compositions may include, for example, detergent formulations for ware-washing applications, detergent compositions for washing of clothes, and laundry additives such as peroxygen bleaches, fabric softeners, enzymes and related products. Pouching cleaning compositions presents the added problem of highly-alkaline contents which can interact with polyvinyl alcohol (PVA) films, which surprisingly severely reduces their solubility, strength, or both. The presence of borate in cleaning

20 compositions (e.g. those containing perborate bleaches) can cause cross-linking of the PVA, reducing its solubility in water. The prior art has attempted to minimize the deleterious effects of borate ions by including a borate scavenger such as sorbitol in the film formulation.

The use of PVA films to contain cleaning compositions is further hampered by variations in solubility caused by the range of water temperatures employed. PVA films of the art generally exhibit varying

30 solubilities in hot (above about 49 °C), warm (about 35 °C) and cold (below about 21 °C) water, depending on the residual acetate content. In addition to the need for rapid film solubility under a variety of wash conditions, the films must be stable over typical storage periods and under a variety of environmental conditions. For example, a film pouch containing a detergent product may be stored under conditions of moderate temperature and humidity, under high temperature and low humidity, or high temperature and

35 high humidity. The latter is not uncommon in certain areas of the Southeastern United States. In high humidity conditions, water can penetrate the film, and if an alkaline detergent is present, can have an adverse impact on the film's integrity. One approach to correcting this problem has been to modify or restrict the amount of alkaline material within the pouch. This can, however, have an adverse impact on the cleaning performance. Another problem with water-soluble PVA film pouches for fabric laundering is the

40 adverse effect of the PVA on cleaning performance.

United States Patent 3,892,905 issued to Albert discloses a cold-water soluble film which may be useful when packaging detergent. Albert, however, does not solve the problem of insolubilization due to alkaline or borate-containing compounds. Great Britain Patent Application 2,090,603, to Sonenstein, describes a

45 packaging film having both hot and cold-water solubility and made from a blend of polyvinyl alcohol and polyacrylic acid. The acrylic acid polymer acts as an alkalinity scavenger, but as the acrylic acids become neutralized, the blend loses its resistance to alkalinity and becomes brittle. The polymers of Sonenstein are not compatible, and preferably are made separately, then blended.

This means an extra process step, and the blend may result in a poor quality film. Dunlop, Jr., US-A-3,198,740 shows a cold-water soluble detergent packet of PVA containing a granular detergent having a

50 hydrated salt to maintain moisture in the film, but without apparent benefit to solubility. United States Patent 4,115,292 issued to Richardson et al shows compositions with enzymes embedded in water-soluble PVA strips, which are in turn encased in a water-soluble film pouch which may be PVA. Lowell et al, US-A-3,005,809 describes copolymers of PVA with 4-10 mole percent of a crotonic acid salt from which films can be made to package neutral, chlorine-liberating compounds. Lowell et al does not teach or suggest any

55 solubility benefits when the films are used to package alkaline or borate-containing detergent compositions.

Inskip, US-A-3,689,469 describes a hot-water soluble copolymer of about 100% hydrolyzed vinyl acetate and about 2 to 6 weight percent methyl methacrylate, and is made to minimize the presence of acid groups. The copolymer can be hydrolyzed using a basic catalyst to form lactone groups, and has utility as

a textile yarn warp-sizing agent. Neher, US-A-2,328,922 and Kenyon, US-A-2,403,004 disclose copolymers of vinyl acetate and acrylic esters, and teach lactone formation to obtain insoluble films. Takigawa, US-A-3,409,598 teaches a process for formation of a water-soluble film using a copolymer of vinyl acetate and an acrylic ester. United States Patents 3,513,142 issued to Blumberg, and 4,155,893 issued to Fujimoto
 5 disclose copolymers of vinyl acetate and a carboxylic ester-containing comonomer. Schulz et al, US-A-4,557,852 describes polymeric sheets which do not include polyvinyl alcohol, but are addition polymers containing high amounts of water-insoluble monomers such as alkyl acrylates and water-soluble anionic monomers such as acrylic salts, and is directed to maintaining flexibility of the sheet during storage. Kaufmann et al, US-A-4,626,372 discloses a PVA film having a polyhydroxy compound which reacts with
 10 borate to afford the film good solubility in the presence of borate. Roulett et al, US-A-4,544,698, describes a PVA and latex combination used as gas-tight moisture resistant coating agents for packaging materials. The latex may include acrylates or methacrylates and vinylidene polychloride polymerized with acrylate, methacrylate or itaconic acid. US-A-4,156,047 discloses water-soluble films formed from a polyethylene glycol plasticizer and a resin mixture containing low and medium molecular weight polyvinyl alcohols and
 15 optionally a copolymer of vinyl alcohol and an ethylenically unsaturated ester.

The problem of enclosing an alkaline or borate-containing laundry product in a water-soluble pouch, which is sufficiently strong for a commercial product, remains storage stable for durations and under environmental conditions typically encountered, and remains water-soluble over a range of wash/rinse temperatures typically encountered in the household, has not been successfully resolved. Accordingly, it is
 20 an object of the present invention to provide a water-soluble film and process for making the same which retains its water solubility in the presence of an alkaline or a borate-containing cleaning composition.

It is another object of the present invention to provide a free-standing film which is water-soluble and stable during storage over a wide range of temperatures and humidities.

It is another object of the present invention to provide a water-soluble film which can be used to
 25 package a cleaning composition and does not have deleterious effects on the performance thereof.

It is yet another object of the present invention to provide a dissolvable laundry additive packet which can be used with alkaline or borate-containing laundry additives.

It is another object to provide a pre-measured, conveniently packaged dose of cleaning composition which is easily stored, handled and delivered to a washing machine, and will rapidly release the cleaning
 30 composition into the wash liquor.

SUMMARY OF THE PRESENT INVENTION

In one embodiment, the present invention is a film formed from a resin having a vinyl acetate monomer
 35 copolymerized with a comonomer selected from a hereinafter defined group. After such copolymerization, and a conversion step, the comonomers are characterized by the presence of an anionic species, and are hereinafter referred to as "nonhydrolyzable" comonomers. The conversion step comprises at least a base catalyzed saponification step, in an organic solvent, to convert residual acetate groups to alcohols, and to produce the anionic species characterizing the nonhydrolyzable comonomer. In some cases, the presence
 40 of adjacent alcohols and carboxylic esters causes the formation of internal lactone rings. By the additional conversion step of subsequently treating the resin with a base, the lactones can also be converted to the anionic form, resulting in an anionic resin from which a film can be made. This latter step is a hydrolysis step. It has been surprisingly found that by selecting the type and content of comonomer, the molecular weight of the PVA resin, and the degrees of hydrolysis of the vinyl acetate, lactonization and ionomer
 45 content, and depending on the type of base used to neutralize the copolymer, a film can be made which exhibits relatively temperature-independent water solubility, and is not rendered insoluble by alkaline or borate-containing detergent compositions. Further, the film is sufficiently strong to be formed into a free-standing pouch which may be used to package cleaning compositions, particularly alkaline or borate-containing cleaning compositions. The film is resistant to insolubilization caused by high humidity storage
 50 conditions, hence is stable over a typical storage shelf life. The films can be produced from a single polymer solution, without the need for making separate polymer solutions, which may be incompatible when mixed for film production. In a second aspect of the present invention, the films are formed into pouches and are used as soluble delivery means for cleaning compositions. Such cleaning compositions include, but are not limited to dry granular, liquid and mulled detergent compositions, bleaches, fabric softeners,
 55 dishwashing detergents, combinations thereof, and other compositions for improving the aesthetics, feel, sanitation or cleanliness of fabrics or wares. The invention is particularly well suited for containing detergent mulls such as those described in European published patent application numbers 0,158,464, filed March 21, 1985, entitled "Low-temperature effective detergent compositions and delivery systems therefor", and

0,234,867, filed February 19, 1987, entitled "Concentrated non-phosphate detergent paste compositions", both of which are assigned to the same assignee as the present invention. These mulls may be highly viscous gels or pastes and include relatively high concentrations of nonionic surfactants for effective removal of oily soils. The mulls are formulated to have alkaline detergent builders which aid in particulate soil removal, and are formulated to provide optimum cleaning power, not for ease of delivery. The preferred delivery method, both for convenience and accuracy, is to include a pre-measured amount of the mull within the water-soluble pouch of the present invention.

It is therefore an advantage of the present invention that high-surfactant, high-builder detergent mulls can be conveniently packaged, stored and delivered.

It is another advantage of the present invention that the films used to package laundry additives remain soluble over the entire range of typical wash temperatures and times.

It is another advantage that the films of the present invention will retain their solubility in contact with alkaline or borate-containing detergents.

It is yet another advantage that the films can be made from a single polymer resin solution.

It is still another advantage of the present invention that the films and film pouches containing detergent remain storage stable over a broad range of environmental conditions.

DETAILED DESCRIPTION OF THE PREFERRED EMBODIMENTS

In a first embodiment, the present invention comprises a free-standing film of a vinyl acetate monomer copolymerized with a comonomer which is converted to yield the nonhydrolyzable comonomer containing an anionic species. Preferably, the anionic species characterizing the nonhydrolyzable comonomer is a carboxylate or sulfonate. Residual acetate groups commonly found in PVA resins are susceptible to alkaline hydrolysis when the resin, or a film made therefrom is exposed to a source of alkalinity. As used herein, the term nonhydrolyzable comonomer is defined to include those repeating units in a PVA copolymer not normally susceptible to hydrolysis by such sources of alkalinity. The nonhydrolyzable comonomers are characterized by the presence of an anionic group, and may be derived from carboxylic acids and salts thereof, carboxylic esters, amides, imides, acyl halides, anhydrides and sulfonates, and impart a degree of water solubility to the resin. This water solubility of the resin should be such that films produced therefrom, having a thickness between 25 to 127 μm (1 to 5 mils), will disperse and substantially dissolve in 70°-130°F (21-54°C) water in less than fifteen minutes, preferably less than five minutes. Subsequent to copolymerization, the nonhydrolyzable comonomer results from the conversion step(s) of, saponification (which also hydrolyzes acetate groups of the polymer to alcohols), or saponification followed by alkaline hydrolysis. The latter hydrolysis step is used when the comonomer is such that lactones are formed as a result of the saponification step. As used herein, the term saponification includes either a base-catalyzed hydrolysis in an organic solvent, or a base catalyzed hydrolysis in an organic solvent followed by the neutralization of excess base and removal of solvent. Preferred bases to catalyze the saponification are the alkali metal hydroxides, including sodium and potassium hydroxide. The organic solvent need not be exclusively organic solvent, but may include some water. Also as used herein, hydrolysis refers to the conversion, usually in a predominately aqueous medium, of a neutral molecule, (e.g. a lactone) to an anionic form, by a source of alkalinity. The presence of adjacent alcohols and carboxylic esters causes internal lactonization of the copolymer resin, but in the presence of a base such as an alkali metal hydroxide, the lactone rings open to form anionic groups, i.e., the salts of the resulting carboxylic acids. Depending on the degree of hydrolysis of the polyvinyl alcohol, the type and percent nonhydrolyzable comonomer content, the degree of lactone conversion, and the type of base used in the hydrolysis step, the resulting resin can be formulated to exhibit varying degrees of water solubility and desired stability characteristics. The anionic groups may be formed during resin or film production, or after film formation. Films can be made with the inventive resin as is known in the art, for example, by solution casting or extrusion, and may be used to pouch gel or mull detergent compositions. Such mulls include detergent builders containing relatively high levels of nonionic surfactants to yield superior oily soil cleaning performance.

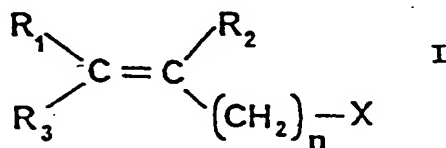
A second embodiment of the present invention comprises a film, made as described in the first embodiment, and fabricated into a pouch. Contained within is a cleaning composition which preferably is a highly viscous, gel or paste detergent composition containing at least one nonionic surfactant and an alkaline builder.

Copolymeric Resins

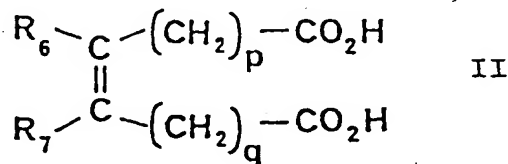
Polyvinyl alcohol (PVA) resin is widely used as a film forming material, and has good strength and water solubility characteristics. Two parameters significantly affecting PVA solubility are molecular weight and degree of hydrolysis. Commercially available films range in weight average molecular weight from 10,000 to 100,000 g/mole. Percent hydrolysis of such commercial PVA films is generally 70% to 100%. Because PVA is made by polymerizing vinyl acetate and subsequently hydrolyzing the resin, PVA can and typically does include residual acetates. The term "polyvinyl alcohol" thus includes vinyl alcohol and vinyl acetate copolymers. For solubility purposes, a high degree of hydrolysis, e.g., 95% renders the film relatively slowly soluble in water. Lower degrees of hydrolysis, e.g. 80-95%, improve solubility rates. In an alkaline environment however, these films become relatively insoluble due to the continued hydrolysis of the partially hydrolyzed film. Higher molecular weight films generally exhibit the best mechanical properties, e.g., impact strength, however solubility rates may be reduced.

It has been surprisingly found that films of the present invention, which are capable of being made into pouches, are storage stable, rapidly soluble over a wide temperature range and are not deleterious to cleaning performance, can be produced from vinyl acetate copolymerized with 2-6 mole percent of a comonomer, to an extent to yield a resin with a molecular weight characterized by a viscosity of between 4 to 35 mPa.s (cPs) as measured in a 4% solution at 25°C, the resin being saponified such that there are 0-10% residual acetate groups, and the comonomers being selected such that subsequent to polymerization, they are converted to nonhydrolyzable comonomers having an anionic charge. As used herein, unless otherwise noted, the resin viscosity is measured after copolymerization and saponification, but before any further treatment of the resin. Mole percentage of comonomer is a measure of the ratio of the number of moles of comonomer to the number of moles of vinyl acetate plus comonomer. In the present invention the resin viscosity should be in the range of between 4-35 mPa.s (cPs), and the mole percentage nonhydrolyzable comonomer is 1-6 percent. Generally, it is desirable to increase the percentage of nonhydrolyzable comonomer as resin viscosity increases, up to the limit of the range. The most preferred nonhydrolyzable comonomer is that which results from the conversion of the methyl acrylate comonomer. The most preferred mole percentage of this nonhydrolyzable anionic comonomer is 3-5%, and it is further most preferred that the resulting resin have a viscosity of 10-20 mPa.s (cPs).

The comonomers which, when copolymerized with vinyl acetate and converted, result in the nonhydrolyzable comonomers having an anionic species are selected from the group of comonomers consisting of unsaturated acids such as acrylic, methacrylic, *cis* 2-butenic, 3-butenic, cinnamic, phenylcinnamic, pentenoic, methylene malonic, the alkali metal and ammonium salts thereof and the acyl halide derivatives thereof; unsaturated esters, amides, and acyl halides of the following structure I:



wherein R_1 , R_2 and R_3 are H, or alkyl, aryl or hydroxyalkyl groups, n is 0 or 1, and X is $-CO_2R_4$, $-C(O)NR_4R_5$ or $-COY$ (wherein R_4 is H, or an alkyl, aryl, alkenyl, hydroxyalkyl, oxyalkyl or cyanoalkyl group, R_5 is H or an alkyl, aryl or hydroxyalkyl group, and Y is a halide); unsaturated diacids and their stereoisomers of the following structure II:



wherein p and q are integers from 0-5, R_6 and R_7 are H, or alkyl or aryl groups, and alkali metal and ammonium salts thereof; anhydrides, acyclic and cyclic esters, amides and imides derived from structure II; unsaturated sulfonic acids and derivatives thereof, and mixtures thereof.

Most suitable comonomers include acrylic acid, methacrylic acid, methylene malonic acid, methyl acrylate, methyl methacrylate, acrylamide, maleic and itaconic acid anhydrides, methyl esters of maleic and itaconic acids, vinyl sulfonate, and mixtures thereof. Conversion of the comonomer to the anionic, nonhydrolyzable comonomer is accomplished by saponification as defined hereinbefore.

5 Some comonomers that are carboxylic acid derivatives, e.g., methyl acrylate and methyl methacrylate, yield lactones on saponification, owing to the presence of adjacent carboxylic esters and alcohols. It has been further surprisingly found that films produced from such lactonized resins do not have acceptable solubility characteristics. For such resins the conversion to anionic form requires alkaline hydrolysis following saponification. The alkaline material used to convert lactones to anionic form may be added
10 before, during or after film production. Operable alkaline materials include but are not limited to alkali metal and alkaline earth metal hydroxides, particularly sodium, lithium and potassium hydroxide, and quaternary ammonium hydroxides, particularly tetraethanol and tetraethyl ammonium hydroxides. Depending on the alkaline material selected, the character of the resulting film can be altered somewhat. For example, solubility of the film is greatest when lithium hydroxide is employed, followed by the sodium, potassium,
15 and quaternary ammonium hydroxides. Film strength is greatest when the quaternary ammonium compounds are used. The alkaline material is added in an amount sufficient to attain the desired mole percentage nonhydrolyzable comonomer, i.e., 1-6 mole percent.

Conversion of the lactone to anionic form may occur as part of the resin or film production process, or after the film has been made but before it is intended to dissolve in water. The introduction of a cleaning
20 composition to the film will result in a degree of anion formation if the cleaning composition is sufficiently alkaline.

Other Film Components

25 The following components are also present in the films of the present invention, and may be added to the resin during film production. A plasticizer is added to the resin to plasticize the copolymeric resin and allow film formation therefrom. Generally any plasticizer known in the art for use with PVA resins will function with the present invention. Preferred are aliphatic polyols, especially ethylene glycol, propylene glycol, glycerol, trimethylolpropane, polyethylene glycol, and mixtures thereof. Particularly preferred is a
30 mixture of polyethylene glycol having a molecular weight of 200-400 g/mole, and glycerol. The total plasticizer content is 0 to 45% by weight of the film composition, preferably 15 to 30 wt % of the film.

A surfactant may be added to the resin mixture to aid in film production by reducing foaming and helping to ensure dispersion and wetting of the composition ingredients. Preferred for this purpose are ethoxylated aliphatic alcohols and ethoxylated alkylphenols. The surfactant may be added in an amount of
35 from 0% to 1.0%, preferably from 0.01% to 0.05%.

To improve solubility of the film in contact with borate-containing additives, a borate scavenger may be added. The borate scavenger is preferably a polyhydroxy compound (PHC) capable of binding to the borate to form a borate-PHC complex. A number of PHC compounds are known in the art to complex with borate such as sorbitol, mannitol, catechol and pentaerythritol. Sorbitol is preferred, and may be added in an
40 amount of from 0 to 30%, preferably from 5 to 20%. A more detailed disclosure of the use of polyhydroxy borate scavengers can be found in US Patent 4,626,372 issued to Kaufmann et al and assigned to the same assignee as the present invention.

Other film additives as known in the art may be included by mixing with the resin. These include antioxidants, release agents, antiblocking agents, and antifoamers, all of which are added in amounts
45 sufficient to perform their intended function as known in the art and generally between 0 and 1% by weight. Film thickness may vary from 25 to 127 μm (1.0 to 5.0 mils), preferably 38 to 64 μm (1.5 to 2.5 mils).

In a second embodiment, the films are used in combination with liquid, solid, granular, paste or mull cleaning compositions to result in a pre-measured, water-soluble packet for cleaning purposes. The cleaning composition may advantageously contain relatively high levels of nonionic surfactants and/or
50 alkaline builders for superior cleaning performance, and/or borate-releasing compounds to provide oxidizing power effective against organic stains. The films of the present invention retain their desired solubility, strength and stability characteristics despite the presence of such alkaline builders or borate, which render ordinary PVA films insoluble, unstable or both. The alkaline cleaning compositions are generally defined as those which generate a pH of greater than 8 when dissolved to a level of 1% in an aqueous medium.
55 Borate-containing cleaning compositions are generally defined as those yielding a borate ion concentration, in water, of greater than about $2.0 \times 10^{-4}\text{M}$. A more detailed description of an example of a detergent mull for which the films of the present invention are particularly adapted for delivering can be found in the previously described European application numbers 0,158,464, and 0,234,867.

The amounts of builders and surfactants which can be included can vary considerably depending on the nature of the builders, the final desired viscosity and the amount of water added to the surfactant system. Other additives commonly found in detergent compositions can be included in the formulations herein. These include but are not limited to additional surfactants, fluorescent whitening agents, oxidants, corrosion inhibiting agents, anti-redeposition agents, enzymes, fabric softeners, perfumes, dyes and pigments. The detergent composition herein may include phosphate or nonphosphate builders.

The following nonlimiting examples are provided to further illustrate the present invention.

Example A

A copolymeric resin was made by copolymerizing vinyl acetate and methyl acrylate to yield about 30 g of the copolymer having a 20,000-25,000 g/mole weight average molecular weight (with an approximate viscosity of 6 mPa.s (cPs)) and 4.5 mole percent methyl acrylate. The resin was saponified to convert 100% of the acetate groups to alcohols and to cause the formation of lactones. The resin had an initial lactone mole percentage of about 4.5%, and a melting temperature of 206 °C. About 30 g of the resin was added to about 190 g of deionized water, and stirred to disperse. About 4 g of a plasticizer, plus about 2 g of a borate scavenger were added to the resin and small quantities (under about 0.5%) of an antiblocking/release agent and an antioxidant were added. The dispersion was heated for about two hours at 60-70 °C to fully dissolve the resin. To this solution sufficient NaOH was added, with heating, to hydrolyze about 1 to 4 mole percent of the lactone groups to anionic form.

The solution was heated for an additional five hours at 60-70 °C to complete the hydrolysis, and was then slowly cooled to about 23 °C and deaerated. The solution was cast on a stainless steel plate using a film applicator with a 0.2 cm clearance. The resulting film was dried at 61 °C for about 30 minutes, cooled to room temperature, and removed from the plate. This procedure yielded a film about 64 µm (2.5 mils) thick, and containing about 70.3% copolymer, 14.3% plasticizer, 7.2% borate scavenger, and 8.2% water.

All of the solubility data were obtained by placing the film in a test device (a 35mm format slide having a 3cm x 4.5cm aperture) in a 600 ml beaker containing about 325 ml of deionized water. Washing machine agitation was simulated by stirring the test solution with a magnetic stirrer at a speed sufficient to result in a vortex extending downward for about 20% of the solution depth. In simulations involving borate, $\text{Na}_2\text{B}_4\text{O}_7$ was added to the water to result in a borate concentration of about $1.7 \times 10^{-3}\text{M}$, and the pH was adjusted to 10.7 with sodium carbonate/bicarbonate. Solubilities of films stored in contact with alkaline cleaning products were determined after the films were removed from contact with the cleaning products and any residual cleaning product adhering to the films was wiped off. Film solubilities were visually evaluated as percentage film residue remaining after 300 seconds in the stirred beaker. Separate studies showed that if the film fully dissolved after 300 seconds in the beaker, no undissolved film residue would be expected from pouched cleaning products in actual use conditions.

Examples B - I

Example B was made as described for Example A, with the copolymeric resin polymerized to have a molecular weight corresponding to about 10 mPa.s (cPs) instead of the 6 mPa.s (cPs). Examples C, D and E were made as described for Example A, but were polymerized to have viscosities of 14 mPa.s (cPs), 17 mPa.s (cPs) and 30 mPa.s (cPs), respectively. Example F was made as Example A with methyl methacrylate instead of methyl acrylate, and with a viscosity of about 15 mPa.s (cPs). Examples G was made by copolymerizing vinyl acetate and maleic anhydride, and had a viscosity of 17 mPa.s (cPs). Example G did not, however, require the subsequent alkaline hydrolysis step of Example A, as the comonomer of Example G was already in anionic form. Examples H and I are prior art polymers of 88% hydrolyzed PVA.

EXPERIMENTAL RESULTS

I. Effects of Resin Viscosity and Copolymer Type and Percent on Alkaline Stability

The alkaline stability of films using various PVA copolymer resins was observed for the following films. Long term film storage in contact with an alkaline detergent was simulated by storing the films in a saturated NaCl solution with the pH adjusted with NaOH to about 12. Dissolution was observed after storage times of 2, 4, 8 and 24 hours in the solution. This test, termed an "accelerated test", simulated in 2 and 4 hours the effect of actual storage for one and two weeks at 32 °C/85% RH. The 8 and 24 hour storage

conditions simulated prolonged actual storage at high humidity. Results are given as percent film remaining after 300 sec in a beaker under the test conditions as outlined previously. Zero percent film remaining indicates desired solubility. The dissolution medium was 21 °C water.

Table 1

Film	Resin Viscosity mPa.s (cPs)	Comonomer		Mole % Residual Acetate	Solubility (% Film residue after 300 sec. in 21 °C water)			
		Type	Total Mole %		2	4	8	24 ⁽¹⁾
A	6	Acrylate*	4.5	0	0	0	0	0
B	10	Acrylate*	4.5	0	0	0	0	0
C	14	Acrylate*	4.5	≤1	0	0	0	0
D	17	Acrylate*	4.5	0	0	0	0	0
E	30	Acrylate*	4.5	0	0	0	0	0
F	15	Methacrylate*	2.7	0	Trace	0	0	0
G	17	Maleate	2.3	3-5	0	0	0	0
H	5	None	-	12	0	50	100	-
I	13	None	-	12	0	100	100	-

*methyl esters

⁽¹⁾Hours in accelerated test solution

This table illustrates that films A-G which are prepared in accordance with the present invention, maintain their solubility under extreme alkaline storage conditions. Films H and I, which are prior art films of vinyl alcohol and vinyl acetate, quickly lose their solubility.

II. Long-term Stability of Film A with Alkaline Paste Detergent

Pouches of an alkaline paste detergent containing a nonionic surfactant, sodium tripolyphosphate, Na₂CO₃, silicate, protease, and a fragrance were prepared using Films D and H. These pouches were exposed to the following storage conditions in a cycling temperature/humidity room, and monitored for film solubility. The cycling room is designed to cycle temperature and humidity from 21 °C/87% RH to 32 °C/65% RH and back over a 24 hour period. These conditions simulate actual weather conditions found in humid regions of the United States.

Table 2

Solubility (21°C water)
% Film Residue After 300 Sec.

Film	Cycling Room 8 weeks	21°C/50% R.H. 6 weeks + Cycling Room 3 weeks
D	0	0
H	80	75

Table 2 demonstrates that the films of the present invention are not insolubilized by hot and/or humid environmental conditions, whereas the prior art PVA film (film H) became, for practical purposes, insoluble under the same conditions.

III. Stability of Film C with Additional Cleaning Products

Table 3

Product	Solubility ⁽¹⁾	
	Film C	Film H
Dry Detergent		
5% Perborate (pH 10.7*)	0	Trace
Dry Bleach		
15% Perborate (pH 11.2*)	0	25%
Dry Automatic		
Dishwashing Detergent (PH 10.3*)	0	10%
* of a 1% solution		
(1) Percent of film remaining after 300 sec. in 21°C water following storage in a cycling room for 4 weeks.		

Table 3 shows the usefulness of the films of the present invention with borate-containing, and highly alkaline additives. It is thought that the anionic nature of the films functions to repel borate anions, and to prevent cross-linking which renders prior art films insoluble.

It has been surprisingly found that molecular weight as represented by viscosity of a 4% polymer solution, and comonomer type and content can impact the cleaning performance of laundry detergents on certain soils, (e.g., on clay soil). Cleaning performance was evaluated by measuring percentage soil removal as a change in fabric reflectance. Swatches of cotton fabric were prepared and stained with BANDY BLACK clay (a trademarked product of the H. C. Spinks Clay Co.), and washed in a commercially available washing machine. Test conditions included 68L of 38°C water at a hardness of 100 ppm (Ca²⁺ and Mg²⁺ in a 3:1 ratio). A 1.8 g piece of film and 53.7 g of paste detergent were used in the evaluation.

Reflectance values of the swatches were measured on a Gardner colorimeter before and after the wash, and the data were analyzed using the Kubelka-Munk equation.

IV. Effect of Polymer Solution Viscosity and Anionic Nonhydrolyzable Comonomer Content on Cleaning Performance

Table 4

Film	Resin Viscosity ⁽¹⁾ mPa.s(cPs)	Mole% Anionic Comonomer	Cleaning Performance (% Soil Removal)
A	6	3.4	92
C	14	3.4	90
E	30	3.4	87
H	5	0	90
I	13	0	80

⁽¹⁾ Measured as a 4% aqueous solution at 25 °C.

It is beneficial, for film strength reasons, to have as high a molecular weight (viscosity) as possible. High molecular weight films of the prior art, however result in poor clay soil performance (a 13 mPa.s (cPs) prior art film yielded about a 10% decrease in cleaning performance over a 5 mPa.s (cPs) prior art film). The films of the present invention, however, show only slight decreases in cleaning performance as viscosity is increased from 6 (film A) to 14 mPa.s (cPs) (film C) and 30 mPa.s (cPs) (film E). For example,

film E of the present invention, at a resin viscosity of 30 mPa.s (cPs), exhibits better cleaning performance than a 13 mPa.s (cPs) film (film I) of the prior art.

V. Effect of Anionic Nonhydrolyzable Comonomer Content on Initial Solubility

Table 5

Film	Solubility (% Film Residue After 300 Sec.)	
	4°C/Water	21°C/Borate
B Anionic	0	0
15 B Neutral	50	25
C Anionic	0	0
C Neutral	50	25
20 D Anionic	0	0
D Neutral	100	100

25 Table 5 shows the neutral copolymer films (e.g. with the comonomer in lactone form) do not dissolve completely in cold or borate-containing water. When the films are in anionic form, i.e., the lactones are converted to the anionic comonomer, however, complete initial dissolution is achieved.

VI. Effect of Anionic Nonhydrolyzable Comonomer Content on Cleaning Performance

30 The degree of anion content in the copolymer films affects the clay-soil removal efficiency of the paste detergent as well as the initial solubility exhibited in the previous example. This effect was demonstrated by controlling the amount of hydrolysis of lactone groups of film D to vary the anion content of the resin. Cleaning performance was measured as described for Table 4, above.

Table 6

	Mole Percent Anionic Nonhydrolyzable Comonomer (Film D)	Percent Soil Removal
	3.4	91
	2.3	90
	1.2	87
45	0	84

Table 6 shows that at a given viscosity level of the films of the present invention, better clay soil removal can be achieved by increasing the anionic content of the film, which can be controlled by the amount of comonomer, and in some cases, by the degree of hydrolysis of intermediate lactone groups.

While described in terms of the presently preferred embodiments, it is to be understood that such disclosure is not to be interpreted as limiting. Various modifications and alterations will no doubt occur to those skilled in the art after having read the above disclosure. Accordingly, it is intended that the appended claims be interpreted as covering all alterations and modifications as fall within the scope of the invention.

Claims

1. A water-soluble polymeric film and additive combination comprising

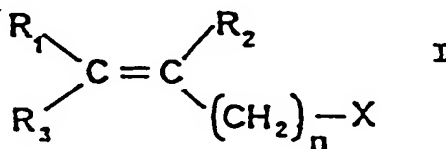
(a) a water-soluble film 25-127 μm (1-5 mils) thick; and

(b) an alkaline or borate-containing additive, at least partially enclosed by the film wherein the film will dissolve when placed in an aqueous medium, freeing the additive;

characterised in that the film is formed from a copolymer resin of vinyl alcohol having 0-10 mole percent residual acetate groups and 1-6 mole percent of a non-hydrolyzable anionic comonomer converted from the group of comonomers consisting of

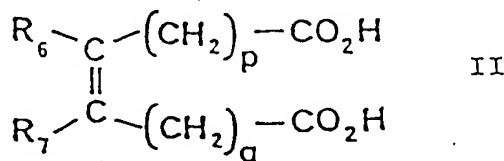
(i) unsaturated acids such as acrylic, methacrylic, cis 2-butenoic, 3-butenoic, cinnamic, phenylcinnamic, pentenoic, methylene malonic, the alkali metal and ammonium salts thereof and the acyl halide derivatives thereof;

(ii) unsaturated esters, amides, and acyl halides of the following structure I:



wherein R_1 , R_2 and R_3 are H, or alkyl, aryl or hydroxyalkyl groups, n is 0 or 1, and X is $-\text{CO}_2R_4$, $-\text{C}(\text{O})\text{NR}_4\text{R}_5$ or $-\text{COY}$, wherein R_4 is H, or an alkyl, aryl, alkenyl, hydroxyalkyl, oxyalkyl or cyanoalkyl group, R_5 is H or an alkyl, aryl or hydroxyalkyl group, and y is a halide;

(iii) unsaturated diacids and their stereoisomers of the following structure II:



wherein p and q are integers from 0-5, R_6 and R_7 are H, or alkyl or aryl groups, and alkali metal and ammonium salts thereof;

(iv) anhydrides, acyclic and cyclic esters, amides and imides derived from structure II;

(v) unsaturated sulfonic acids and derivatives thereof; and

(vi) mixtures thereof,

the resin being polymerized to an extent to result in a resin viscosity, when dissolved in 25 °C water to a level of 4% of between 4-35 mPa.s (cPs), the film including a plasticizing-effective amount of a plasticizer.

2. The combination of claim 1, characterised in that the copolymeric resin is produced by copolymerizing vinyl acetate with 2-6 mole percent of a comonomer selected from comonomers (i)-(vi) to yield a copolymer resin, subsequently saponifying the resin to hydrolyse 90-100 mole percent acetate groups to alcohols, and converting sufficient of the comonomer to result in 1-6 mole percent of the nonhydrolyzable anionic comonomer.

3. The combination of claim 1 or claim 2 wherein the comonomer is selected from the group consisting of acrylic acid, methacrylic acid, methylene malonic acid, methyl acrylate, methyl methacrylate, acrylamide, maleic and itaconic acid anhydrides, methyl esters of maleic and itaconic acids, vinyl sulfonate, and mixtures thereof.

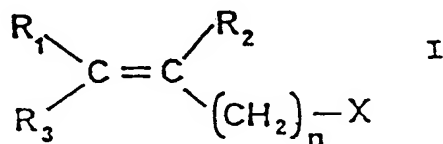
4. The combination of claim 2 or claim 3 wherein the conversion of the comonomer to the non-hydrolyzable comonomer occurs during the saponification.

5. The combination of claim 2 or 3 wherein the conversion of the comonomer to the nonhydrolyzable comonomer further includes a hydrolysis with an alkaline material, following the saponification.

6. The combination of claim 5 wherein the alkaline material is an alkaline-earth metal, alkali-metal, or quaternary ammonium hydroxide, and mixtures thereof.

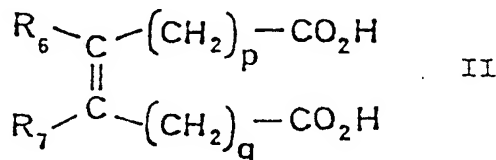
7. The combination of claim 5 or claim 6 wherein the alkaline material is included in the additive.
8. The combination of any claims 1-7 and further including up to 30 weight percent of a borate scavenger in the film.

9. A method for introducing an additive to an aqueous solution comprising
- (a) enclosing an additive in a sealed water-soluble copolymeric film material produced by copolymerizing vinyl acetate with 2-6 mole% of a comonomer to yield a copolymer resin, subsequently saponifying the resin to hydrolyze 90-100 mole percent acetate groups to alcohols, the copolymer resin characterized by a viscosity, when dissolved to a level of 4% in 25°C water, of between 4 to 35 mPa.s (cPs), and converting sufficient of the comonomer to result in 1-6 mole percent of a nonhydrolyzable comonomer having an anionic charge, wherein the comonomer is selected from the group consisting of
- (i) unsaturated acids such as acrylic, methacrylic, cis 2-butenic, 3-butenic, cinnamic, phenylcinnamic, pentenoic, methylene malonic, the alkali metal and ammonium salts thereof and the acyl halide derivatives thereof;
- (ii) unsaturated esters, amides, and acyl halides of the following structure I:



wherein R_1 , R_2 and R_3 are H, or alkyl, aryl or hydroxyalkyl groups, n is 0 or 1, and X is $-CO_2R_4$, $-C(O)NR_4R_5$ or $-COY$, wherein R_4 is H, or an alkyl, aryl, alkenyl, hydroxyalkyl, oxyalkyl or cyanoalkyl group, R_5 is H or an alkyl, aryl or hydroxyalkyl group, and Y is a halide;

(iii) unsaturated diacids and their stereoisomers of the following structure II:



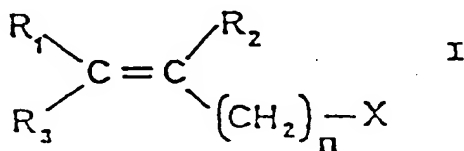
wherein p and q are integers from 0-5, R_6 and R_7 are H, or alkyl or aryl groups, and alkali metal and ammonium salts thereof;

- (iv) anhydrides, acyclic and cyclic esters, amides and imides derived from structure II;
- (v) unsaturated sulfonic acids and derivatives thereof; and
- (vi) mixtures thereof; and
- (b) contacting the copolymeric material plus additive with an aqueous solution for a sufficient time to dissolve the polymeric material and disperse the additive contained therein.
10. The method of claim 9 wherein the comonomer is selected from the group consisting of acrylic acid, methacrylic acid, methylene malonic acid, methyl acrylate, methyl methacrylate, acrylamide, maleic and itaconic acid anhydrides, methyl esters of maleic and itaconic acids, vinyl sulfonate, and mixtures thereof.
11. The method of claim 9 wherein the conversion of the comonomer to the nonhydrolyzable comonomer occurs during the saponification.
12. The method of claim 9 wherein the conversion of the comonomer to the nonhydrolyzable comonomer further includes a hydrolysis with an alkaline material, following the saponification.
13. The method of claim 12 wherein the alkaline material is an alkaline-earth metal, alkali-metal, or quaternary ammonium hydroxide, and mixtures thereof.

14. The method of claim 12 wherein the alkaline material is included in the additive.
15. The method of claim 9 and further including adding up to 30 weight percent of a borate scavenger in the film.
16. The method of claim 9 wherein the film has a thickness of 25-127 μm (1-5 mils).
17. The method of claim 9 wherein the film includes a plasticising-effective amount of a plasticiser.

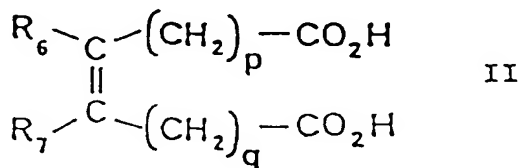
10 Patentansprüche

1. Kombination aus einem wasserlöslichen polymeren Film und einem Zusatzstoff, umfassend
- (a) einen 25-127 μm (1-5 mils) dicken wasserlöslichen Film; und
- (b) einen alkalischen oder Borat-enthaltenden Zusatzstoff, der mindestens teilweise von dem Film umhüllt ist, wobei sich der Film, wenn er in wäßriges Medium gegeben wird, auflöst und den Zusatzstoff freisetzt;
- dadurch **gekennzeichnet**, daß der Film aus einem Copolymerharz aus einem Vinylalkohol mit 0-10 Mol-% Restacetatgruppen und 1-6 Mol-% eines nicht-hydrolysisierbaren anionischen Comonomeren, umgewandelt aus der Gruppe aus Comonomeren, die besteht aus
- (i) ungesättigten Säuren, wie Acrylsäure, Methacrylsäure, cis-But-2-ensäure, But-3-ensäure, Zimtsäure, Phenylzimtsäure, Pentensäure, Methylenmalonsäure, den Alkalimetall- und Ammoniumsalzen davon und den Acylhalogenidderivaten davon;
- (ii) ungesättigten Estern, Amiden und Acylhalogeniden der folgenden Struktur I:



worin R_1 , R_2 und R_3 H oder Alkyl-, Aryl- oder Hydroxyalkylgruppen bedeuten, n 0 oder 1 bedeutet und X $-\text{CO}_2R_4$, $-\text{C}(\text{O})\text{NR}_4R_5$ oder $-\text{COY}$ bedeutet, worin R_4 für H oder eine Alkyl-, Aryl-, Alkenyl-, Hydroxyalkyl-, Oxyalkyl- oder Cyanoalkylgruppe steht, R_5 für H oder eine Alkyl-, Aryl- oder Hydroxyalkylgruppe steht und Y für ein Halogenid steht;

(iii) ungesättigten Disäuren und ihren Stereoisomeren der folgenden Struktur II:



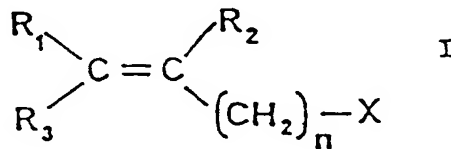
worin p und q ganze Zahlen von 0-5 bedeuten, R_6 und R_7 H oder Alkyl- oder Arylgruppen bedeuten, und den Alkalimetall- und Ammoniumsalzen davon;

(iv) Anhydriden, acyclischen und cyclischen Estern, Amiden und Imiden, die sich von der Struktur II ableiten;

(v) ungesättigten Sulfonsäuren und Derivaten davon; und

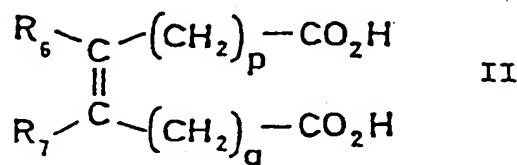
(vi) den Gemischen davon gebildet worden ist, wobei das Harz in dem Umfang polymerisiert worden ist, daß eine Harzviskosität nach dem Auflösen in 25 °C Wasser bei einem Gehalt von 4% zwischen 4-35 mPa.s (cPs) erhalten wird und der Film eine weichmachende Menge eines Weichmachers enthält.

2. Kombination nach Anspruch 1, dadurch **gekennzeichnet**, daß das Copolymerharz durch Copolymerisation von Vinylacetat mit 2-6 Mol-% eines Comonomeren, ausgewählt aus den Comonomeren (i) - (vi), gebildet worden ist, wobei das Harz anschließend verseift wird, um 90-100 Mol-% der Acetatgruppen zu Alkoholen zu hydrolysieren, und um ausreichend Comonomeres umzuwandeln, daß 1-6 Mol-% der nicht-hydrolysierbaren anionischen Comonomeren gebildet werden.
3. Kombination nach Anspruch 1 oder 2, dadurch **gekennzeichnet**, daß das Comonomere ausgewählt wird aus der Gruppe, die besteht aus Acrylsäure, Methacrylsäure, Methylenmalonsäure, Methylacrylat, Methylmethacrylat, Acrylamid, Malein- und Itaconsäureanhydriden, Methylestern von Malein- und Itaconsäuren, Vinylsulfonat und Gemischen davon.
4. Kombination nach Anspruch 2 oder 3, dadurch **gekennzeichnet**, daß die Umwandlung des Comonomeren in das nicht-hydrolysierbare Comonomere während der Verseifung stattfindet.
5. Kombination nach Anspruch 2 oder 3, dadurch **gekennzeichnet**, daß die Umwandlung des Comonomeren in das nicht-hydrolysierbare Comonomere weiter eine Hydrolyse mit einem alkalischen Material, gefolgt von der Verseifung, umfaßt.
6. Kombination nach Anspruch 5, dadurch **gekennzeichnet**, daß das alkalische Material ein Erdalkalimetall-, ein Alkalimetall- oder ein quaternäres Ammoniumhydroxid und Gemische davon ist.
7. Kombination nach Anspruch 5 oder 6, dadurch **gekennzeichnet**, daß das alkalische Material in dem Zusatzstoff enthalten ist.
8. Kombination nach irgendeinem der Ansprüche 1 bis 7, dadurch **gekennzeichnet**, daß es zusätzlich bis zu 30 Gew.-% eines Boratreinigungsmittels in dem Film enthält.
9. Verfahren zur Einführung eines Zusatzstoffes in eine wäßrige Lösung, dadurch **gekennzeichnet**, daß
 (a) ein Zusatzstoff mit einem versiegelten wasserlöslichen Copolymerfilmmaterial umhüllt wird, wobei das Filmmaterial durch Copolymerisation von Vinylacetat mit 2-6 Mol-% eines Comonomeren unter Bildung eines Copolymerharzes, anschließend Verseifen des Harzes, um 90-100 Mol-% Acetatgruppen in Alkohol zu hydrolysieren, hergestellt worden ist, das Copolymerharz durch eine Viskosität nach dem Auflösen bis zu einem Gehalt von 4% in 25°C Wasser zwischen 4 und 35 mpa.s (cPs) charakterisiert ist und ausreichend Comonomeres umgewandelt wurde, um 1-6 Mol-% nicht-hydrolysierbares Comonomeres mit anionischer Ladung zu ergeben, wobei das Comonomere aus der Gruppe ausgewählt wurde, die besteht aus
 (i) ungesättigten Säuren, wie Acrylsäure, Methacrylsäure, cis-But-2-ensäure, But-3-ensäure, Zimtsäure, Phenylzimtsäure, Pentensäure, Methylenmalonsäure, den Alkalimetall- und Ammoniumsalzen davon und den Acylhalogenidderivaten davon;
 (ii) ungesättigten Estern, Amiden und Acylhalogeniden der folgenden Struktur I:



worin R_1 , R_2 und R_3 H oder Alkyl-, Aryl- oder Hydroxyalkylgruppen bedeuten, n 0 oder 1 bedeutet und X $-CO_2R_4$, $-C(O)NR_4R_5$ oder $-COY$ bedeutet, worin R_4 für H oder eine Alkyl-, Aryl-, Alkenyl-, Hydroxyalkyl-, Oxyalkyl- oder Cyanoalkylgruppe steht, R_5 für H oder eine Alkyl-, Aryl- oder Hydroxyalkylgruppe steht und Y für ein Halogenid steht;

(iii) ungesättigten Disäuren und ihren Stereoisomeren der folgenden Struktur II:



worin p und q ganze Zahlen von 0-5 bedeuten, R₆ und R₇ H oder Alkyl- oder Arylgruppen bedeuten, und den Alkalimetall- und Ammoniumsalzen davon;

(iv) Anhydriden, acyclischen und cyclischen Estern, Amiden und Imiden, die sich von der Struktur II ableiten;

(v) ungesättigten Sulfonsäuren und Derivaten davon; und

(vi) den Gemischen davon; und

(b) das Copolymermaterial plus der Zusatzstoff mit einer wäßrigen Lösung während einer Zeit behandelt werden, die ausreicht, das polymere Material zu lösen und den darin enthaltenen Zusatzstoff zu dispergieren.

10. Verfahren nach Anspruch 9, dadurch **gekennzeichnet**, daß das Comonomere ausgewählt wird aus der Gruppe, die besteht aus Acrylsäure, Methacrylsäure, Methylenmalonsäure, Methylacrylat, Methylmethacrylat, Acrylamid, Malein- und Itaconsäureanhydriden, Methylestern von Malein- und Itaconsäuren, Vinylsulfonat, und Gemischen davon.

11. Verfahren nach Anspruch 9, dadurch **gekennzeichnet**, daß die Umwandlung des Comonomeren in das nicht-hydrolysierbare Comonomere während der Verseifung stattfindet.

12. Verfahren nach Anspruch 9, dadurch **gekennzeichnet**, daß die Umwandlung des Comonomeren in das nicht-hydrolysierbare Comonomere zusätzlich die Hydrolyse mit einem alkalischen Material, gefolgt von der Verseifung, umfaßt.

13. Verfahren nach Anspruch 12, dadurch **gekennzeichnet**, daß das alkalische Material ein Erdaalkalimetall-, Alkalimetall- oder quaternäres Ammoniumhydroxid oder ein Gemisch davon ist.

14. Verfahren nach Anspruch 12, dadurch **gekennzeichnet**, daß das alkalische Material in dem Zusatzstoff enthalten ist.

15. Verfahren nach Anspruch 9, dadurch **gekennzeichnet**, daß es zusätzlich die Zugabe von bis zu 30 Gew.-% Boratreinigungsmittel zu dem Film umfaßt.

16. Verfahren nach Anspruch 9, dadurch **gekennzeichnet**, daß der Film eine Dicke von etwa 25-127 µm (1-5 mils) besitzt.

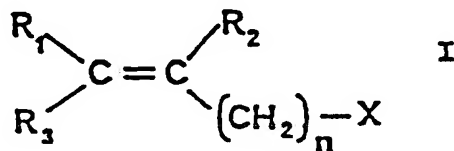
17. Verfahren nach Anspruch 9, dadurch **gekennzeichnet**, daß der Film eine weichmachende Menge eines Weichmachers umfaßt.

Revendications

1. Une combinaison d'une pellicule polymère soluble dans l'eau et d'un additif comprenant
 - (a) une pellicule soluble dans l'eau épaisse d'environ 25-127 µm (1-5 mils) ; et
 - (b) un additif alcalin ou contenant un borate, au moins partiellement enfermé par la pellicule, où la pellicule se dissout lorsqu'elle est placée dans un milieu aqueux pour libérer l'additif ;
 caractérisée en ce que la pellicule est formée à partir d'une résine copolymère d'alcool vinylique ayant 0 à 10 % molaires de groupes acétates résiduels et 1 à 6 % molaires d'un comonomère anionique non hydrolysable obtenu par conversion à partir du groupe de comonomères constitué de :

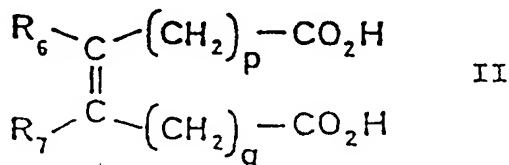
(i) les acides insaturés comme les acides acrylique, méthacrylique, cis-2-buténoïque, 3-buténoïque, cinnamique, phénylcinnamique, penténoïque, méthylènemalonique et leurs sels de métal alcalin et d'ammonium et les halogénures d'acyle qui en dérivent ;

(ii) les esters, amides et halogénures d'acyle insaturés de structure I suivante :



dans laquelle R_1 , R_2 et R_3 sont H ou des groupes alkyles, aryles ou hydroxyalkyles, n est 0 ou 1, et X est $-CO_2R_4$, $-C(O)NR_4R_5$ ou $-COY$, où R_4 est H ou un groupe alkyle, aryle, alcényle, hydroxyalkyle, oxyalkyle ou cyanoalkyle, R_5 est H ou un groupe alkyle, aryle ou hydroxyalkyle, et Y est un halogénure ;

(iii) les diacides insaturés et leurs stéréo-isomères de structure II suivante :



dans laquelle p et q sont des entiers de 0 à 5, R_6 et R_7 sont H ou des groupes alkyles ou aryles et leurs sels de métal alcalin et d'ammonium ;

(iv) les anhydrides, esters acycliques et cycliques, amides et imides dérivés de la structure II ;

(v) les acides sulfoniques insaturés et leurs dérivés ; et

(vi) leurs mélanges,

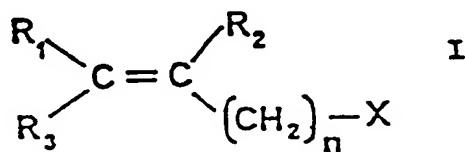
la résine étant polymérisée dans une mesure qui établit une viscosité de la résine, lorsqu'elle est dissoute dans l'eau à 25°C, à la concentration de 4 % entre 4 et 35 mPa.s (cP), la pellicule contenant une quantité efficace pour la plastification d'un plastifiant.

2. La combinaison de la revendication 1, caractérisée en ce que la résine copolymère est produite par copolymérisation d'acétate de vinyle avec 2 à 6 % molaires d'un comonomère choisi parmi les comonomères (i)-(vi), pour former une résine copolymère, puis saponification de la résine pour hydrolyser 90 à 100 % molaires des groupes acétates en alcools et conversion suffisante du comonomère pour obtenir 1 à 6 % molaires du comonomère anionique non hydrolysable.
3. La combinaison de la revendication 1 ou de la revendication 2, dans laquelle le comonomère est choisi dans le groupe constitué par l'acide acrylique, l'acide méthacrylique, l'acide méthylènemalonique, l'acrylate de méthyle, le méthacrylate de méthyle, l'acrylamide, les anhydrides des acides maléique et itaconique, les esters méthyliques des acides maléique et itaconique, le vinylsulfonate et leurs mélanges.
4. La combinaison de la revendication 2 ou de la revendication 3, dans laquelle la conversion du comonomère en le comonomère non hydrolysable a lieu lors de la saponification.
5. La combinaison de la revendication 2 ou 3, dans laquelle la conversion du comonomère en le comonomère non hydrolysable comprend de plus une hydrolyse avec une matière alcaline, puis la saponification.
6. La combinaison de la revendication 5, dans laquelle la matière alcaline est un hydroxyde de métal alcalino-terreux, de métal alcalin ou d'ammonium quaternaire et leurs mélanges.
7. La combinaison de la revendication 5 ou de la revendication 6, dans laquelle la matière alcaline est incluse dans l'additif.

8. La combinaison de l'une quelconque des revendications 1 à 6 comprenant de plus jusqu'à 30 % en poids d'un épurateur de borate dans la pellicule.

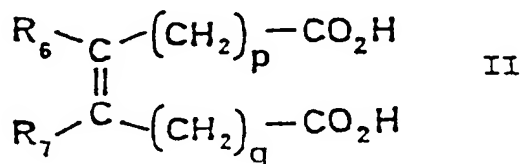
9. Un procédé pour introduire un additif dans une solution aqueuse consistant à :

- (a) enfermer un additif dans une pellicule copolymère étanche soluble dans l'eau, produite par copolymérisation d'acétate de vinyle avec 2 à 6 % molaires d'un comonomère pour former une résine copolymère, puis saponifier la résine pour hydrolyser 90 à 100 % molaires des groupes acétates en alcools, la résine copolymère étant caractérisée par une viscosité, lorsqu'elle est dissoute à la concentration de 4 % dans de l'eau à 25°C, entre 4 et 35 mPa.s (cP), et effectuer la conversion de suffisamment du comonomère pour obtenir 1 à 6 % molaires d'un comonomère non hydrolysable ayant une charge anionique, où le comonomère est choisi dans le groupe constitué de
- (i) les acides insaturés comme les acides acrylique, méthacrylique, cis-2-buténoïque, 3-buténoïque, cinnamique, phénylcinnamique, penténoïque, méthylènemalonique et leurs sels de métal alcalin et d'ammonium et les halogénures d'acyle qui en dérivent ;
- (ii) les esters, amides et halogénures d'acyle insaturés de structure I suivante :



dans laquelle R_1 , R_2 et R_3 sont H ou des groupes alkyles, aryles ou hydroxyalkyles, n est 0 ou 1, et X est $-CO_2R_4$, $-C(O)NR_4R_5$ ou $-COY$, où R_4 est H ou un groupe alkyle, aryle, alcényle, hydroxyalkyle, oxyalkyle ou cyanoalkyle, R_5 est H ou un groupe alkyle, aryle ou hydroxyalkyle, et Y est un halogénure ;

(iii) les diacides insaturés et leurs stéréo-isomères de structure II suivante :



dans laquelle p et q sont des entiers de 0 à 5, R_6 et R_7 sont H ou des groupes alkyles ou aryles et leurs sels de métal alcalin et d'ammonium ;

- (iv) les anhydrides, esters acycliques et cycliques, amides et imides dérivés de la structure II ;
- (v) les acides sulfoniques insaturés et leurs dérivés ; et
- (vi) leurs mélanges ; et
- (b) mettre la matière copolymère plus l'additif en contact avec une solution aqueuse pendant un temps suffisant pour dissoudre la matière polymère et disperser l'additif qui y est contenu.

10. Le procédé de la revendication 9, dans lequel le comonomère est choisi dans le groupe constitué par l'acide acrylique, l'acide méthacrylique, l'acide méthylènemalonique, l'acrylate de méthyle, le méthacrylate de méthyle, l'acrylamide, les anhydrides des acides maléique et itaconique, les esters méthyliques des acides maléique et itaconique, le vinylsulfonate et leurs mélanges.

11. Le procédé de la revendication 9, dans lequel la conversion du comonomère en le comonomère non hydrolysable se produit lors de la saponification.

12. Le procédé de la revendication 9, dans lequel la conversion du comonomère en le comonomère non hydrolysable comprend de plus une hydrolyse avec une matière alcaline, puis la saponification.

13. Le procédé de la revendication 12, dans lequel la matière alcaline est un hydroxyde de métal alcalino-terreux, de métal alcalin ou d'ammonium quaternaire, et leurs mélanges.

14. Le procédé de la revendication 12, dans lequel la matière alcaline est incluse dans l'additif.
15. Le procédé de la revendication 9 comprenant de plus l'addition de jusqu'à 30 % en poids d'un épurateur de borate dans la pellicule.
16. Le procédé de la revendication 9, dans lequel la pellicule a une épaisseur de 25 à 127 μm (1 à 5 mils).
17. Le procédé de la revendication 9, dans lequel la pellicule comprend une quantité efficace pour la plastification d'un plastifiant.